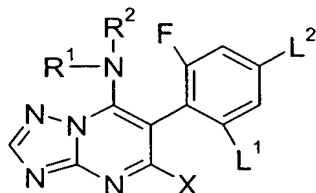


6-(2-Fluorophenyl)-triazolopyrimidines, method for producing them, their use for controlling parasitic fungi and agents containing the same

Abstract

5

6-(2-Fluorophenyl)-triazolopyrimidines of the formula I



in which the substituents are as defined below:

10 R¹ is C₄-C₈-alkyl, C₄-C₈-haloalkyl, substituted C₃-C₈-Cycloalkyl, C₃-C₈-halocycloalkyl, C₅-C₈-alkenyl, C₂-C₈-haloalkenyl, C₃-C₆-cycloalkenyl, C₃-C₆-halocycloalkenyl, C₂-C₈-alkynyl, C₂-C₈-haloalkynyl or phenyl, naphthyl, or a five- or six-membered saturated, partially unsaturated or aromatic heterocycle which contains one to four heteroatoms from the group consisting of O, N and S,

15

R² is hydrogen, C₁-C₃-alkyl or one of the groups mentioned under R¹,

R¹ and R² together with the nitrogen atom to which they are attached may also form a five- or six-membered heterocyclyl or heteroaryl which is attached via N and contain one to three further heteroatoms from the group consisting of O, N and S as ring member,

20

except for piperidin-1-yl optionally substituted by methyl groups;

25 R¹ and/or R² may be substituted according to the description;

L¹ is chlorine or fluorine;

L² is hydrogen,
30 is, if L¹ is fluorine, also fluorine;

X is alkyl;

processes for preparing these compounds, compositions comprising them and their
35 use for controlling phytopathogenic harmful fungi.